abcam

Product datasheet

Anti-Matriptase 2 antibody - Cytoplasmic domain ab56180

★★★★★ 2 Abreviews 6 References 画像数 1

製品の概要

製品名 Anti-Matriptase 2 antibody - Cytoplasmic domain

製品の詳細 Rabbit polyclonal to Matriptase 2 - Cytoplasmic domain

由来種 Rabbit

特異性 ab56180 reacts with Matripase 2 (TMPRSS6) and does not react with the other TMPRSSs.

ab56180 recognizes all of the listed forms of Matripase 2.

 アプリケーション
 適用あり: WB

 種交差性
 交差種: Human

免疫原 Synthetic peptide based on the aminoterminal end (cytoplasmic domain) of Matripase 2 (Human).

特記事項

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

バッファー pH: 7.40

Preservative: 0.05% Sodium azide

Constituents: 50% Glycerol, 2.9% Sodium chloride, PBS

精製度 Immunogen affinity purified

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

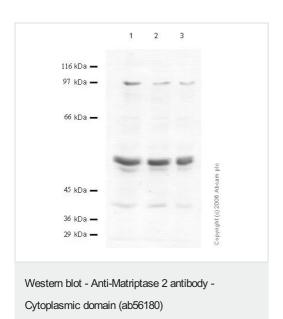
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
WB	★★★★ (1)	1/1000 - 1/5000. Predicted molecular weight: 89 kDa.

ターゲット情報

機能	Serine protease which hydrolyzes a range of proteins including type I collagen, fibronectin and fibrinogen. Can also activate urokinase-type plasminogen activator with low efficiency. May play a specialized role in matrix remodeling processes in liver. Required to sense iron deficiency. Overexpression suppresses activation of the HAMP promoter.
組織特異性	Liver specific.
関連疾患	Defects in TMPRSS6 are the cause of iron-refractory iron deficiency anemia (IRIDA) [MIM:206200]; also known as hypochromic microcytic anemia with defect in iron metabolism or hereditary iron-handling disorder or pseudo-iron-deficiency anemia. Key features include congenital hypochromic microcytic anemia, very low mean corpuscular erythrocyte volume, low transferrin saturation, abnormal iron absorption characterized by no hematologic improvement following treatment with oral iron, and abnormal iron utilization characterized by a sluggish, incomplete response to parenteral iron.
配列類似性	Belongs to the peptidase S1 family. Contains 2 CUB domains. Contains 3 LDL-receptor class A domains. Contains 1 peptidase S1 domain.
ドメイン	Cytoplasmic domain mediates HAMP suppression via proximal promoter element(s).
細胞内局在	Cell membrane.

画像



All lanes : Anti-Matriptase 2 antibody - Cytoplasmic domain (ab56180) at 1/1000 dilution

All lanes: Cell media from human embryonic kidney HEK293 cells

Predicted band size: 89 kDa

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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